erial No. 10/567,113 Docket No. 1004398.002US (4874-7001)

## LISTING OF CLAIMS

This listing of claims replaces all prior versions and listings of claims in the application:

1-27. (Cancelled)

28. (New) A method of treating diseases caused by disorders of nitrergic system and/or dopaminergic system of an organism comprising administering an active ingredient having normalizing effect with respect to nitrergic and dopaminergic systems, wherein the active ingredient is present in a pharmaceutically-acceptable carrier in an amount sufficient for effecting said systems, said active ingredient being a cyclic bioisostere of derivatives of a purine system having a general structural formula

H—CH<sub>2</sub>ONa

R<sup>1</sup> is selected from the group consisting of -H, -NH<sub>2</sub>, -Br, -Cl, -OH, and -COOH;

where R is selected from the group consisting of HO

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A is selected from the group consisting of -N= and -C=;

B is selected from the group consisting of -N= and -C=;

Z is selected from the group consisting of -C= and -N=;

wherein when A is -N=, then B is -N= and Z is -C=,

or pharmacologically acceptable salts thereof.

29. (New) The method as claimed in claim 28, wherein said active ingredient is selected from the group consisting of:

sodium salt of 7-(P-D-ribofuranosile)pyrido[2,3-d]-6H-pyridazine-5,8-dione;

sodium salt of 4-amino-7-(P-D-ribofuranosile)pyrido[2,3-d]-6H-pyridazine-5,8-dione;

sodium salt of 3-bromo-7-(P-D-ribofuranosile)pyrido[2,3-d]-6H-pyridazine-5,8-dione;

disodium salt of 4-hydroxy-7-(P-D-ribofuranosile)pyrido[2,3-d]-6H-pyridazine-5,8-dione;

disodium salt of 3-carboxy-7-(P-D-ribofuranosile)pyrido[2,3-d]-6H-pyridazine-5,8-dione;,

lithium salt of pyrido [2,3-d]-6H-pyridazine-5,8-dione;

sodium salt of pyrido [2,3-d]-6H-pyridazine-5,8-dione;

potassium salt of pyrido [2,3-d]-6H-pyridazine-5,8-dione;

sodium salt of 2-(P-D-ribofuranosile)benzo[d]-3H-pyridazine-l,4-dione;

sodium salt of 5-amino-2-(p-D-ribofuranosile)benzo[d]-3H-pyridazine-l,4-dione;

sodium salt of 6-amino-2-(3-D-ribofuranosile)benzo[d]-3H-pyridazine-1,4-dione;

sodium salt of 5-chloro-2-(P-D-ribofuranosile)benzo[d]-3H-pyridazine-1,4-dione;

disodium salt of 5-hydroxy-2-(P-D-ribofuranosile)benzo[d]-3H-pyridazine-1,4-dione;

lithium salt of 5-amino-benzo[d]-3H-pyridazine-1,4-dione;

sodium salt of 5-amino-benzo[d]-3H-pyridazine-l,4-dione;

potassium salt of 6-amino-benzo[d]-3H-pyridazine-1,4-dione;

disodium salt of 5-hydroxy-benzo[d]-3H-pyridazine-l,4-dione;

disodium salt of 6-carboxy-benzo[d]-3H-pyridazine-1,4-dione;

sodium salt of 7-(P-D-ribofuranosile)pyrazine[2,3-d]-6H-pyridazine-5,8-dione;

sodium salt of 2-amino-7-(P-D-ribofuranosile)pyrazine[2,3-d]-6H-pyridazine-5,8-dione: sodium salt of 3-amino-7-(P-D-ribofuranosile)pyrazine[2,3-d]-6H-pyridazine-5,8-dione; sodium salt of 3-bromo-7-(P-D-ribofuranosile)pvrazine[2,3-d]-6H-pvridazine-5,8-dione; disodium salt of 2-hydroxy-7-(p-D-ribofuranosile)pyrazine[2,3-d]-6H-pyridazine-5,8-dione; disodium salt of 2-carboxy-7-(P-D-ribofuranosile)pyrazine[2,3-d]-6H-pyridazine-5,8-dione; lithium salt of pyrazine[2,3-d]-6H-pyridazine-5,8-dione; sodium salt of pyrazine[2,3-d]-6H-pyridazine-5,8-dione; potassium salt of 3-bromo-pyrazine[2,3-d]-6H- pyridazine-5,8-dione; sodium salt of 2-amino-pyrazine[2,3-d]-6H-pyridazine-5,8-dione; sodium salt of 7-(B-D-ribofuranosile)pyrimido[4,5-d]-6H-pyridazine-5,8-dione; sodium salt of 2-amino-7-(β-D-ribofuranosile)pyrimido[4,5-d]-6H-pyridazine-5,8-dione; sodium salt of 4-amino-7-(β-D-ribofuranosile)pyrimido[4,5-d]-6H-pyridazine-5,8-dione; sodium salt of 2-bromo-7-(β-D-ribofuranosile)pyrimido[4,5-d]-6H-pyridazine-5,8-dione; sodium salt of 4-hydroxy-7-(β-D-ribofuranosile)pyrimido[4,5-d]-6H-pyridazine-5,8-dione; sodium salt of 4-carboxy-7-(B-D-ribofuranosile)pyrimido[4,5-d]-6H-pyridazine-5,8-dione; lithium salt of pyrimido[4,5-d]-6H-pyridazine-5,8-dione; sodium salt of 2-amino-pyrimido[4,5-d]-6H-pyridazine-5,8-dione; and potassium salt of 4-bromo-pyrimido[4,5-d]-6H-pyridazine -5,8-dione.

30. (New) The method as claimed in claim 28, wherein the active ingredient is used as a neuroprotector in a pharmaceutical composition for protection of a nervous system.

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Response to Office Action dated January 8, 2009

31. (New) The method as claimed in claim 28, wherein the active ingredient is used

in a pharmaceutical composition for improvement of cognitive function and normalization of

psychophysiological status.

32. (New) The method as claimed in claim 28, wherein the active ingredient is used

in a pharmaceutical composition of anxiolytic and antidepressive action.

33. (New) The method as claimed in claim 28, wherein the diseases are selected

from the group consisting of: diseases caused by drug abuse; insomnia; sexual disorders/sexual

dysfunction; gastro-intestinal disorders; psychoses; inorganic psychoses; affective disorders;

personality disorders; psychiatric disorders of mood; schizophrenia and schizoaffective

disorders; polydipsia; bipolar disorders; dysphoric mania; anxiety and associated diseases; obesity; bacterial infections of the central nervous system; disorders of learning; disorders of

memory; Parkinson's disease; neurodegenerative diseases; Alzheimer's disease; depression;

extrapyramidal side effects of neuroleptics; hypothalamic-pituitary effects; vascular and

 $cardiovas cular\ diseases;\ dystonia;\ dyskinesia;\ hyperkinesis;\ dementia;\ is chemia;\ motion$ 

disorders; hypertension; and diseases caused by a hyperactive immune system.

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